## PROTOCOL REVISIONS

<table>
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| 2                | 02nd June 2015 | Text reformatted and clarified, information updated Questionnaires created and updated | 1                            | **General updates/reformatting**  
Clarification of secondary endpoints (obstetric/neonatal)  
**Additional clinical information to be collected, to allow sensitivity analyses in future individual patient data meta-analyses**  
Internal pilot phase interviews refined and made specific  
Inclusion/exclusion criteria updated  
Clarification of withdrawal  
Pessary criteria updated  
Ultrasound accreditation method updated  
Qualitative interviews/questionnaires and Health Economics questionnaires included and information updated  
PAG information updated  
Study monitoring procedures updated  
Trial Steering Committee details updated  
Data Monitoring Committee details updated  
Secondary end point definitions added (Appendix 5) |
| 3                | 5th August 2015 | Change of measurement which defines "short cervix" and confers eligibility for randomisation | 2                            | Change of measurement which defines a short cervix from ≤ 30mm to ≤ 35mm. The anticipated population centile (≤ 30th centile) is unchanged.                                                                                 |
| 4                | 10th January 2018 | Clarification of text, modification of the data collection and safety                | 5                            | • Clarification of Co-enrolment  
• Modification of data collection to include hospitals, which are not participating STOPPIT2 sites  
• Safety reporting guidance modified for clarity.  
• Increase in the number of participants from 1850 to 2500 for cervical length scanning to facilitate randomisation which remains at 500.                                      |